



Europäisches Patentamt
European Patent Office
Office européen des brevets



Publication number: **0 529 693 A2**

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: **92119095.5**

(51) Int. Cl. 5: **C09D 5/14**

(22) Date of filing: **12.10.89**

This application was filed on 07 - 11 - 1992 as a divisional application to the application mentioned under INID code 60.

(30) Priority: **13.10.88 GB 8824003**
12.05.89 GB 8910970

(43) Date of publication of application:
03.03.93 Bulletin 93/09

(60) Publication number of the earlier application in accordance with Art. 76 EPC: **0 364 271**

(84) Designated Contracting States:
DE ES FR GB GR IT NL SE

(71) Applicant: **COURTAULDS COATINGS (HOLDINGS) LIMITED**
50 George Street
London W1A 2BB(GB)

(72) Inventor: **Hunter, Julian Edward**
Meadow Rise, Kenton
Newcastle-Upon-Tyne NE5 4SW(GB)
Inventor: **Reid, James**
150C Whitehall Road
Gateshead, Tyne and Wear(GB)
Inventor: **Arnold, David Edward John**
132 Moorside North
Fenham, Newcastle-Upon-Tyne NE4 9DY(GB)
Inventor: **Hails, George**
4 Granby Close
Sunnyside, Newcastle-Upon-Tyne NE16 5ND(GB)
Inventor: **Baxter, Kenneth Ford**
5 Greybourne Gardens
Sunderland(GB)
Inventor: **Andrews, Adrian Ferguson**
22 Shaws Park
Hexham, Northumberland(GB)
Inventor: **Nunn, Michael John**
Low Walden Farmhouse, Low Walden
Nr. Hexham, Northumberland(GB)

(74) Representative: **Hale, Stephen Geoffrey et al**
J.Y. & G.W. Johnson Furnival House 14/18
High Holborn
London WC1V 6DE (GB)

(54) **Antifouling coatings.**

(57) Fouling of a ship's hull is inhibited by applying to the hull a coating composition having marine biocidal properties and comprising as binder an acid-functional film-forming polymer whose acid groups are blocked by hydrolysable blocking groups which are monoamine groups forming organic-solvent-soluble amine salts of the polymer.

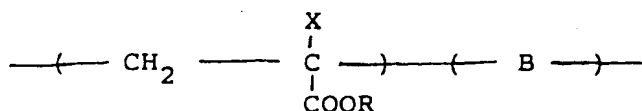
EP 0 529 693 A2

This invention relates to antifouling coatings. An antifouling coating is used as a top coat on ships' hulls to inhibit the settlement and growth of marine organisms such as barnacles and algae, generally by release of a biocide for the marine organisms.

Traditionally, antifouling coatings have comprised a relatively inert binder with a biocidal pigment which is leached from the paint. Among the binders which have been used are vinyl resins, particularly a vinyl chloride/vinyl acetate copolymer, and rosin. The vinyl resins are seawater-insoluble and paints based on them use a high pigment concentration so that there is contact between pigment particles to ensure leaching. Rosin is a hard brittle resin which is slightly soluble in seawater. The biocidal pigment is very gradually leached out of the matrix of rosin binder in use, leaving a skeletal matrix of rosin which becomes washed off the hull surface to allow leaching of the biocidal pigment from deep within the paint film.

The most successful antifouling paints in recent years have been self-polishing copolymer paints based on a polymeric binder to which biocidal triorganotin moieties are chemically bound and from which the biocidal moieties are gradually hydrolysed by seawater, as described for example in GB-A-1457590. The polymer from which the triorganotin moieties have been hydrolysed becomes soluble in seawater, so that as the outermost paint layer becomes depleted of biocide it is swept off the surface of the hull by the movement of the ship through seawater. Self-polishing copolymer paints which release non-biocidal moieties are described in EP-B-69559 and EP-A-232006.

WO84/02915 describes an antifouling paint having a hydrolysable film-forming water-insoluble seawater-erodible polymeric binder having recurring groups represented by the formula:



where X is hydrogen or methyl, R is an alkyl, aryl, aralkyl or triorganosilyl moiety and B is the residue of an ethylenically unsaturated comonomer. It has been found in practice that the less readily hydrolysable groups R such as benzyl, aminoalkyl or haloalkyl groups do not give a polymer which dissolves in seawater, whereas the more readily hydrolysable groups R such as trialkylsilyl groups give a polymer which rapidly hydrolyses to a mechanically weak film in seawater.

JP-A-54-64633 describes a marine antifouling biocide which is a long-chain (12 to 18 carbon atoms) linear aliphatic primary amine or salt thereof. JP-A-54-110322 describes certain long-chain (12 to 18 carbon atoms) linear aliphatic secondary and tertiary amines as marine antifouling agents.

US-A-4675051 describes a marine antifouling paint which is gradually dissolved in seawater and which comprises a binder which is a resin produced by the reaction of rosin and an aliphatic polyamine containing at least one primary or secondary amine group.

An antifouling coating composition according to the invention comprises a pigment and as binder an acid-functional film-forming polymer whose acid groups are blocked by hydrolysable blocking groups, the composition including an ingredient having marine biocidal properties, and is characterised in that the hydrolysable blocking group is a monoamine group which forms an organic-solvent-soluble amine salt of the polymer.

The invention further provides a process for inhibiting fouling of a ship's hull, comprising applying to the hull a coating composition having marine biocidal properties and comprising as binder an acid-functional film-forming polymer whose acid groups are blocked by hydrolysable blocking groups, characterised in that the hydrolysable blocking group is a monoamine group which forms an organic-solvent-soluble amine salt of the polymer.

The acid-functional polymer is preferably a carboxylic-acid-functional polymer of equivalent weight 240 to 600. A preferred acid-functional polymer is an addition copolymer of one or more olefinically unsaturated acids or anhydrides, for example acrylic acid, methacrylic acid, maleic acid, maleic anhydride, fumaric acid, itaconic acid or itaconic anhydride, vinyl benzoic acid (for example p-vinyl benzoic acid), 3-butenic acid or beta-carboxy-ethyl acrylate or methacrylate, with at least one olefinically unsaturated comonomer. Copolymers of methacrylic acid or acrylic acid are preferred. (The preferred equivalent weight of 240 to 600 corresponds to an acrylic acid content of 14.3 to 35.8% by weight and a methacrylic acid content of 16.7 to 41.7% by weight.) The acid monomer is preferably copolymerised with one or more comonomers which are unreactive with acid groups, for example acrylic or methacrylic esters such as methyl acrylate, methyl methacrylate, ethyl acrylate, butyl acrylate or 2-ethylhexyl methacrylate, styrene, acrylonitrile, vinyl acetate, vinyl butyrate, vinyl chloride, or vinyl pyridine. Terpolymers may be preferred, for example methyl

methacrylate or ethyl methacrylate which tend to form a hard film can be used in conjunction with an acrylate such as ethyl acrylate or particularly an alkyl acrylate of 3 to 8 carbon atoms in the alkyl moiety such as butyl acrylate which helps to form a more flexible film. A substituted acrylate ester can be used as one of the comonomers, for example. Such an acid polymer preferably has a molecular weight of 1,000 to 100,000. The equivalent weight of the acid polymer (calculated as acid groups) is most preferably 300 to 440, equivalent to an acrylic acid or methacrylic acid content of about 15 to 30% by weight.

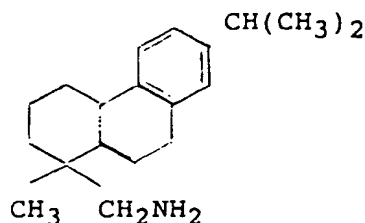
Alternative acid-functional polymers are polymers containing sulphonic acid, phosphonic acid or phosphoric acid (acid phosphate) groups. If alternative acid groups are used they are also preferably present in an addition polymer, for example an addition copolymer of an olefinically unsaturated phosphonic, phosphoric or sulphonic acid. Examples of such unsaturated acids are vinyl phosphonic acid, styrene phosphonic acid, 2-acrylamidopropane phosphonic acid, ethylidene-1,1-diphosphonic acid, hydroxyethyl acrylate monophosphate, vinyl sulphonic acid, 2-acrylamido-2-methylpropane sulphonic acid, methallyl sulphonic acid and styrene sulphonic acid. Polymers containing stronger acid groups such as sulphonic acid groups may have a higher equivalent weight for example in the range 500 to 5000, preferably 1000 to 2000.

The monoamine which is used to form the amine salt of the acid-functional polymer preferably includes at least one organic group containing at least 8 carbon atoms, more preferably 8 to 20 carbon atoms, and is preferably an amine which is toxic to marine organisms. If such an amine is used the resulting amine salt can be a clear antifouling varnish or can be pigmented. The monoamine can for example be a diterpene-derived amine of the formula:



where R^1 is a monovalent hydrocarbon group derived from a diterpene and R^2 and R^3 are each independently hydrogen, an alkyl group having 1 to 18 carbon atoms or an aryl group having 6 to 12 carbon atoms.

An amine of the formula (I) acts as a marine antifouling biocide. The amine is preferably derived from rosin. The main constituent of rosin is abietic acid, which is mixed with other diterpene acids. The amine is preferably a primary or secondary amine. Secondary amines, for example those in which R^2 is a methyl group, may be the more effective biocides against fouling by animals such as barnacles, whereas primary amines may be the more effective biocides against algae. A primary amine derived from rosin is dehydroabietylamine sold commercially as "Rosin Amine D". Its main constituent is:



A corresponding secondary or tertiary amine, for example an N-methyl or N,N-dimethyl derivative of Rosin Amine D, can alternatively be used.

The amine of formula (I) can be used as a mixture with one or more other high molecular weight amines in forming a binder for an antifouling paint according to the invention. It can for example be used with a long-chain (12 to 20 carbon atoms) aliphatic amine such as dodecyl amine, hexadecyl amine, octadecyl amine or oleyl amine, or a mixture of such amines, for example those sold as tallow amine, hydrogenated tallow amine, coconut amine, or N-methyl coconut amine. Such a mixture of amines preferably contains at least 50% by weight of the amine of formula (I), for example 60 to 90%.

The toxic amine can alternatively be an aliphatic amine containing an organic group of 12 to 20 carbon atoms, for example a straight-chain alkyl or alkenyl amine such as dodecyl amine, hexadecyl amine, octadecyl amine or oleyl amine or mixtures of amines derived from aliphatic groups present in natural fats

and oils such as tallow amine or hydrogenated tallow amine or coconut amine. Alternative amines which can be used as the blocking group are aralkylamines such as those sold commercially as "phenalkamines".

If a non-biocidal amine is used to form the amine salt of the acid-functional polymer the coating composition should contain a marine biocide. The coating preferably contains a pigment, which may be the same as the marine biocide.

The amine salt formed from the acid-functional polymer is substantially insoluble in water and controls the rate of dissolution of the acid-functional polymer in seawater. The amine salt gradually dissociates on prolonged immersion in seawater, for example on a ship's hull in service. The amine is gradually released into the seawater. The remaining acid-functional polymer is gradually converted to free acid or anion form and becomes seawater-soluble and is gradually swept from the hull of the ship. The paints containing the amine salts of an acid-functional polymer thus act as self-polishing coatings. When a biocidal amine is used, the paints have properties very similar to known organotin copolymer paints, releasing polymer-bound biocide with the polymer binder itself gradually becoming smoothly dissolved from the ship's hull in service.

The blocked acid-functional polymer can be prepared by addition polymerisation of the corresponding blocked acid-functional monomer, i.e. an amine salt of a polymerisable ethylenically unsaturated acid such as acrylic or methacrylic acid, with one or more comonomers. Polymerisation is preferably carried out in an organic solvent such as xylene, toluene, butyl acetate, butanol, butoxyethanol or methoxypropyl acetate at a temperature of 60 to 100 °C using a free radical catalyst such as benzoyl peroxide or azobisisobutyronitrile. The amine salt is preferably formed in solution in a polar organic solvent by reaction of an acidic monomer such as acrylic or methacrylic acid with the amine to produce an amine salt and polymerised without isolating the salt, although it can be isolated if desired. The blocked acid-functional polymer can alternatively be prepared by reacting an acid-functional copolymer having free carboxyl groups with an amine as blocking agent to form an amine salt. The amine salt can be formed by simply mixing the amine and a solution of the acid-functional polymer, preferably in an organic solvent such as an aromatic hydrocarbon, a ketone, an alcohol or an ether alcohol.

The amine-blocked acid-functional polymer can be mixed with pigment using conventional paint blending procedures to provide a composition having a pigment volume concentration of, for example, 25 to 55%. The pigment is preferably a sparingly soluble pigment having a solubility in seawater of from 0.5 to 10 parts per million by weight, for example cuprous oxide, cuprous thiocyanate, zinc oxide, zinc ethylene bis(dithiocarbamate), zinc dimethyl dithiocarbamate, zinc diethyl dithiocarbamate or cuprous ethylene bis(dithiocarbamate). These sparingly soluble pigments which are copper and zinc compounds are generally marine biocides. These pigments produce water-soluble metal compounds on reaction with seawater so that the pigment particles do not survive at the paint surface. Mixtures of sparingly soluble pigments can be used, for example cuprous oxide, cuprous thiocyanate or zinc ethylene bis(dithiocarbamate), which are highly effective biocidal pigments, can be mixed with zinc oxide, which is less effective as a biocide but dissolves slightly more rapidly in seawater. Both the amine of formula (I) and the amine-blocked acid-functional polymer can be mixed with a basic pigment such as cuprous oxide or zinc oxide without gelation of the binder, unlike acid-functional polymers containing free carboxylic acid groups. The amine salt protects the acid groups against gelation by a basic pigment.

The paint composition can additionally or alternatively contain a pigment which is not reactive with seawater and may be highly insoluble in seawater (solubility below 0.5 part per million by weight) such as titanium dioxide or ferric oxide or an organic pigment such as phthalocyanine pigment. Such highly insoluble pigments are preferably used at less than 40% by weight of the total pigment component of the paint, most preferably less than 20%.

The antifouling paint can also contain a non-metalliferous biocide for marine organisms, for example tetramethyl thiuram disulphide, methylene bis(thiocyanate), captan, a substituted isothiazolone or 2-methylthio-4-t-butylamino-6-cyclopropylamino-s-triazine.

The antifouling coating composition of the invention is generally applied from a solution in an organic solvent, for example an aromatic hydrocarbon such as xylene or toluene, an aliphatic hydrocarbon such as white spirit, an ester such as butyl acetate, ethoxyethyl acetate or methoxypropyl acetate, an alcohol such as butanol or butoxy-ethanol or a ketone such as methyl isobutyl ketone or methyl isoamyl ketone.

Alternatively, the anti fouling composition of the invention can be an aqueous composition containing water and a water-miscible cosolvent. Examples of cosolvents which can be used are alcohols such as butanol, glycol ethers such as methoxypropanol, methoxyethanol, butoxyethanol and ethoxyethanol and esters thereof such as methoxypropyl acetate. The acid-functional polymers blocked with an amine containing an organic group of at least 8 carbon atoms are soluble in such mixtures of water and cosolvent but are substantially insoluble in water.

The invention is illustrated by the following Examples.

Example 1

A 40% solution of a 24/56/20 copolymer of methacrylic acid/ethyl methacrylate/methoxy ethyl acrylate was prepared by solution polymerisation using a free radical catalyst in a 1:1 by volume mixture of xylene and butanol. Rosin amine D was added to the polymer solution in an amount of 1.03 amine groups per acid group in the polymer.

The antifouling properties of the varnish of Example 1 were tested using a leaching test. In this test a plaque coated with a film of the varnish of Example 1 was immersed in a tank of synthetic seawater and was removed for a day once a week and immersed in a smaller tank of seawater. The seawater from the smaller tank was then tested each time for toxicity against *Artemia* (brine shrimp) and *Amphora* (unicellular algae) marine organisms.

The seawater samples which had been in contact with the coating of Example 1 showed positive and substantially constant toxicity over the 8 week test period indicating that marine biocide continued to be leached from the paint over a prolonged period.

By comparison, when the acid copolymer solution was applied as a coating without reacting with an amine and was tested, the leached samples were seen to be non-toxic.

Examples 2 and 3

7.5 g of the polymer amine salt solution of Example 1 was mixed with 1.5 g cuprous oxide (Example 2) and 1.5 g zinc oxide (Example 3) to form antifouling paints.

The viscosity of the paints of Examples 2 and 3 was monitored over a hundred hours after mixing by an ICI cone and plate viscometer at 25 °C. The paint of Example 2 stayed at a stable viscosity of less than 10 poise over the 100 hour test. The viscosity of the paint of Example 3 was less than 40 poise at the end of the 100 hour test. By comparison, paints formed by mixing the acid-functional polymer with cuprous oxide or zinc oxide without reacting the polymer with the amine showed a rise in viscosity to over 100 poise (far too high for spray application) within 80 hours.

Example 4

85% by volume of the polymer amine salt solution of Example 1 was milled with 14.4% by volume cuprous oxide, 0.25% by volume bentonite and 0.35% by volume silica aerogel to form an antifouling paint.

The paint was tested in a rotor disc test of the type described in GB-A-1457590 and showed a gradual decrease in thickness over 60 days' immersion. The polishing rate was similar to that of a commercial triorganotin copolymer antifouling paint. By comparison, when the acid copolymer was mixed with the cuprous oxide pigment without being first reacted with the amine the resulting paint was removed from the rotor disc after one day's immersion.

Example 5

A 30/20/50 copolymer of acrylic acid/methyl methacrylate/butyl acrylate was prepared in xylene/butanol solution as described in Example 1. Rosin Amine D was added to the copolymer in an amount equivalent to the acid groups in the polymer. The resulting solution was milled with pigments, plasticiser and structuring agents to form a paint containing, by volume, 30% copolymer amine salt, 18% cuprous oxide, 1.3% zinc oxide, 4% tricresyl phosphate and 2% structuring agents.

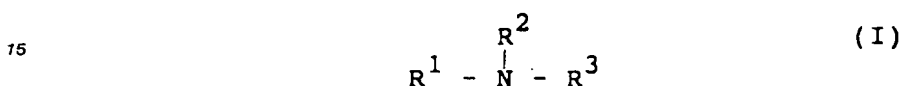
The paint was sprayed on a plaque which was attached to a metal plate which was immersed in the sea at a site off the south coast of England rich in fouling organisms. The painted plaque showed substantially no fouling after 12 months' immersion.

The paint was tested in a rotor disc test and showed a gradual depletion in thickness at a rate substantially the same as that of a commercial self-polishing copolymer antifouling paint.

Claims

1. An antifouling coating composition comprising a pigment and as a binder an acid-functional film-forming polymer whose acid groups are blocked by hydrolysable blocking groups, the composition including an ingredient having marine biocidal properties, characterised in that the hydrolysable blocking group is a monoamine group which forms an organic-solvent-soluble amine salt of the polymer.

2. An antifouling coating composition according to claim 1, characterised in that the acid-functional polymer is a carboxylic-acid-functional polymer of equivalent weight 240 to 600.
- 5 3. An antifouling coating composition according to claim 1 or 2, characterised in that the monoamine providing the amine group includes at least one organic group containing 8 to 20 carbon atoms.
4. An antifouling coating composition according to any of claims 1 to 3, characterised in that the monoamine has marine biocidal properties.
- 10 5. An antifouling coating composition according to claim 3, characterised in that the monoamine providing the amine group is an amine of the formula



where R¹ is a monovalent hydrocarbon group derived from a diterpene and R² and R³ are each independently hydrogen, an alkyl group having 1-18 carbon atoms or an aryl group having 6-12 carbon atoms,

6. An antifouling coating composition according to any of any of claims 1 to 3, characterised in that the monoamine is non-biocidal and the coating composition contains a marine biocide.
- 25 7. An antifouling coating composition according to claim 6, characterised in that the pigment is a marine biocide.
8. An antifouling coating composition according to any of claims 1 to 7, characterised in that the pigment is a copper or zinc compound which is sparingly soluble in seawater.
- 30 9. A process for inhibiting fouling of a ship's hull, comprising applying to the hull a coating composition having marine biocidal properties and comprising as binder an acid-functional film-forming polymer whose acid groups are blocked by hydrolysable blocking groups, characterised in that the hydrolysable blocking group is a monoamine group which forms an organic-solvent-soluble amine salt of the polymer.
- 35